CLAIMS

- 1. Nucleotide sequence encoding a polypeptide capable of interacting with topoisomerase III α .
- 2. Nucleotide sequence according to
 Claim 1, characterized in that it encodes a polypeptide
 comprising all or part of the polypeptide sequence
 SEQ ID No. 4 or of its derivatives.
- 3. Nucleotide sequence according to either of Claims 1 and 2, characterized in that it comprises all or part of the nucleotide sequence SEQ ID No. 3 or of its derivatives.
 - 4. Polypeptide, characterized in that it is capable of interacting with topoisomerase $\text{III}\alpha$.
- 5. Polypeptide according to Claim 4, characterized in that it comprises all or part of the polypeptide sequence SEQ ID No. 4 or of a sequence derived therefrom.
- 6. Polypeptide according to Claim 4,

 20 characterized in that it consists of the polypeptide sequence corresponding to residues 318-662 of the sequence SEQ ID No. 5 or of a sequence derived therefrom.
- 7. Use of a polypeptide or of a polypeptide 25 fragment according to Claims 4 to 6, for slowing down, inhibiting, stimulating or modulating the activity of topoisomerase IIIα.

- 8. Antibody or antibody fragment directed against a polypeptide according to one of Claims 4 to 6.
- 9. Method for detecting or identifying
 5 compounds capable of binding to a polypeptide as
 defined according to one of Claims 4 to 6,
 characterized in that the following steps are carried
 out:
- a a molecule or a mixture containing

 various molecules, optionally unidentified, is brought into contact with a polypeptide as defined according to one of Claims 4 to 6 under conditions allowing the interaction between said polypeptide and said molecule in the case where the latter might possess affinity for said polypeptide, and,
 - b the molecules bound to said polypeptide are detected and/or isolated.
- 10. Method for detecting or identifying compounds capable of modulating or inhibiting the interaction between topoisomerase III-α and a polypeptide as defined according to Claims 4 to 6, characterized in that the following steps are carried out:
- a the binding of topoisomerase III α or of a 25 fragment thereof to said polypeptide is carried out;

- b a compound to be tested for its capacity to inhibit the binding between topoisomerase III α and said polypeptide is added;
- c the displacement or inhibition of the binding of topoisomerase III α to said polypeptide is determined:
 - d the compounds which prevent or which impede the binding between topoisomerase III α and said polypeptide are detected and/or isolated.
- 11. Ligand for a polypeptide as defined according to Claims 4 to 6, capable of being obtained according to the method of Claim 9.
- 12. Ligand capable of modulating or inhibiting the interaction between topoisomerase IIIα
 5 and a polypeptide as defined according to Claims 4 to 6, capable of being obtained according to the method of Claim 10.
- 13. Use of a ligand according to Claim 11 or 12 for the preparation of a medicament intended for the 20 prevention, improvement or treatment of diseases involving a cell cycle dysfunction.
 - 14. Pharmaceutical composition comprising, as active ingredient, at least ligand according to either of Claims 11 and 12 or an antibody according to Claim 8.

- 15. Composition according to Claim 14, intended for modulating slowing down or inhibiting the activity of topoisomerase III α .
- 16. Composition according to either of

 5 Claims 14 and 15, intended for the prevention,
 improvement or treatment of diseases involving a cell
 cycle dysfunction.